

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
29 August 2002 (29.08.2002)

PCT

(10) International Publication Number
WO 02/066404 A1

- (51) International Patent Classification⁷: **C07C 2/32**, B01J 31/22
- (21) International Application Number: PCT/NL01/00149
- (22) International Filing Date: 22 February 2001 (22.02.2001)
- (25) Filing Language: English
- (26) Publication Language: English
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- (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: CATALYST SYSTEM FOR THE TRIMERISATION OF OLEFINS

(57) Abstract: The invention relates to a catalyst system for the selective trimerisation of olefins, which system is based on a titanium complex of formula $(\text{Cp}-\text{B}(\text{R})_n\text{Ar})\text{TiR}^1_3$, wherein: Cp is a cyclopentadienyl type ligand, optionally substituted, B is a bridging group, based on a single atom selected from the groups 13 to 16 inclusive of the Periodic System, Ar is a aromatic group, optionally substituted, R is, independently, hydrogen, or a hydrocarbon residue, optionally being substituted and optionally containing heteroatoms, or groups R and B are joined together to form a ring, n is an integer equal to the (valency of B minus 2), and R^1 is a mono-anionic group, and further comprises an activator. The present catalyst system obviates the use of toxic chromium compounds.

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Title: Catalyst system for the trimerisation of olefins.

The invention relates to a catalyst system for the selective trimerisation of olefins, which system comprises a transition metal complex.

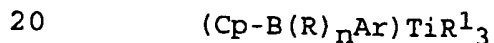
Such a catalyst for trimerisation of olefins is known from
5 EP-A-0608447 and consists of a combination of a metal source, a pyrrole containing compound and a metal alkyl in an electron donor solvent. The transition metal source consists of a chromium, nickel, cobalt, or iron compound, preferably a chromium compound is used.

Because chromium compounds are highly toxic, and therefore need
10 special handling precautions, a catalyst system for the trimerisation of olefins, which is not based on a chromium compound is needed.

A catalyst system has now been found which is not based on a chromium compound, but still shows a high selectivity in the trimerisation of olefins with respect to the trimerisation product.

15 More specifically, the invention relates to a catalyst system as indicated above, which is characterized in that said catalyst comprises

a) a half-sandwich substituted cyclopentadienyl titanium complex of formula



wherein

Cp is a cyclopentadienyl type ligand, optionally substituted,

B is a bridging group, based on a single atom selected from the groups 13 to 16 inclusive of the Periodic System,

25 Ar is a aromatic group, optionally substituted,

R is, independently, hydrogen, or a hydrocarbon residue, optionally being substituted and optionally containing heteroatoms, or groups R and B are joined together to form a ring,

30 n is an integer equal to the (valency of B minus 2), and

R¹ is a mono-anionic group, and

b) an activator.

It is observed that the catalyst system as disclosed in
EP-A-0608447 is preferably a chromium catalyst, but a catalytic system
35 based on a titanium compound, more specifically $\text{TiO}(\text{acac})_2$, was also

tested as a catalyst for the trimerisation of ethylene. The selectivity for hexene-1 was in that case nevertheless rather low. A high selectivity for hexene-1 is industrially very important because of the use of hexene-1 as starting material for the preparation of 5 different kinds of (co)polymers.

It is further observed that trimerisation is in the above-mentioned reference and in the present disclosure defined as the combination of one or more kinds of olefins, wherein the number of olefin, i.e. double, bonds is reduced by two. The term "trimerisation" 10 is thus intended to include "co-trimerisation". So, for example, the number of olefin bonds in the combination of three ethylene units is reduced by two, to one olefin bond, in 1-hexene.

A half-sandwich substituted cyclopentadienyl titanium complex as a catalyst, in the presence of a co-catalyst, is known for example 15 from Macromol. 1999, 32, 4491-4493. This titanium complex does nevertheless not have a bridging group in its structure; moreover, the catalyst system is used for the synthesis of polyethylenes containing significant amounts of butyl branches, and is thus used in a polymerisation process.

20 As is known from EP-A-0 780 353, the properties of a polymer do not change markedly with the addition or removal of one or a few repeating units, contrary to the properties of a product obtained by oligomerisation or trimerisation. A polymerisation catalyst thus results in completely different products than a trimerisation catalyst 25 does.

Half-sandwich cyclopentadienyl titanium complexes of formula $(CpB(R)_2Ar)TiMe_3$ and $(CpB(R)_2Ar)TiCl_3$ are known per se from J. Saßmannshausen et al., J. Organomet. Chem. 1999, 592, 84-94. In these known complexes $B(R)_2Ar$ can be CMe_2Ph , $CHPh_2$ or $SiMe_2Ph$. These 30 complexes were only used as polymerization catalysts; there is no indication at all that these known catalysts could effectively be used for the selective trimerisation of olefins. On the contrary: it is said that the effect of a comparatively weakly coordinated pendant ligand, such as phenyl, on the behaviour of a polymerisation catalyst 35 is difficult to predict. Moreover, a possible favourable effect of a catalyst having a bridging group only consisting of a single atom, for a trimerisation process of ethene to obtain hexene-1, is not mentioned or suggested in this reference.

As mentioned before, Cp is a cyclopentadienyl type ligand, 40 which is optionally substituted.

More preferably, Cp is a cyclopentadienyl, indenyl or fluorenyl group, which may be substituted or not with one to five (cyclopentadienyl), one to seven (indenyl) or one to nine (fluoranyl) substituent alkyl or silyl groups, especially methyl or trimethylsilyl groups.

In the catalyst system according to the invention Ar is an aromatic group, which is optionally substituted; examples thereof are phenyl, naphthalene, anthracene or phenanthrene. This enumeration is not to be regarded as limitative; other aromatic groups can also be used, provided that a coordination complex, based on π -electrons of said group, together with titanium can be formed.

In a preferred embodiment of the invention, the catalyst system comprises a complex of the above given formula, wherein said group B consists of carbon or silicon,
Ar is phenyl, optionally substituted or being part of a larger aromatic entity,

R^1 is a halide, or mono-anionic hydrocarbon residue optionally containing heteroatoms, and
n is 2, then R is a mono-anionic hydrocarbon residue, optionally containing heteroatoms, or
n is 1, then R is a di-anionic hydrocarbon residue, optionally containing heteroatoms.

More preferably the catalyst system of the invention comprises a titanium complex of the above given formula, wherein
Ar is a phenyl group, substituted or not at the meta-or para-position(s),
B is based on a carbon atom,
n is 2, then groups R are, independently, methyl, or ethyl; or
n is 1, then group R is $=CH_2$, or forms when R is C_4H_8 or C_5H_{10} together with group B a dianionic cyclic group,
Cp is C_5H_4 or $C_5H_3(SiMe_3)$, and
 R^1 is chlorine, methyl, or benzyl.

The half-sandwich, substituted cyclopentadienyl titanium complex, forming a part of the present catalyst system, is in a preferred embodiment supported by a carrier. This carrier consists expediently either of a metal oxide, which is selected from the group consisting of alumina, boria, magnesia, thorium, zirconia, silica, or mixtures thereof, or it consists of a polymeric material.

As indicated above, the present catalyst system comprises an activator. Said activator is preferably methylalumoxane, a salt of a non-coordinating anion, or a Lewis acid capable of abstracting an anion from said transition metal complex and thus generating a cationic transition metal species with a non-coordinating anion.

An example of a salt of a non-coordinating anion is N,N-dimethylanilinium tetrakis(pentafluorophenyl)borate, while such a Lewis acid is for example $B(C_6F_5)_3$. It is in this respect observed that any activator can be used provided that it is able to generate a cationic transition metal species with a non-coordinating anion. The term "non-coordinating anion" is meant to indicate the anionic part or derivative of the activator, which not or only weakly coordinates to the cationic form of the present catalyst.

Preferably the activator is methylalumoxane (also known as MAO). The molar ratio of Ti:Al is expediently from 1:100 to 1:1000.

The present catalyst system can further also comprise a scavenger. Examples of a scavenger are $i\text{-Bu}_3\text{Al}$ and $(i\text{-Bu}_2\text{Al})_2\text{O}$. A scavenger is normally used to scavenge impurities from the polymerisation medium to obtain a high productivity.

The invention further relates to a process to trimerize olefinic compounds which comprises carrying out said trimerisation in the presence of a catalyst system, as described above, under trimerisation conditions. Such a trimerisation also comprises co-trimerisation according to the definition given before.

The olefin to be trimerized is preferably selected from $C_2\text{-}C_{20}$ olefins or mixtures of two or more of these olefins. The preferred olefins are ethylene and 1-butene, more preferably ethylene.

The temperature is preferably in the range of from 20-150°C, at a pressure in the range of from 1,5 to 3 MPa.

The invention will further be explained in the following examples.

Experimental section

General considerations

All experiments were performed under a nitrogen atmosphere using standard Schlenk and glovebox techniques. Deuterated solvents (Aldrich, Acros) were dried over Na/K alloy and vacuum transferred before use. Cyclooctane (Aldrich, used as internal standard) was distilled from Na prior to use. Toluene (Aldrich, anhydrous, 99,8%)

was passed over columns of Al_2O_3 (Fluka), BASF R3-11 supported Cu oxygen and molecular sieves (Aldrich, 4Å). Diethyl ether and THF (Aldrich) were dried over Al_2O_3 (Fluka) and the other solvents (Aldrich) were dried over molecular sieves (Aldrich, 4Å). Ethene (AGA 5 polymer grade) was passed over BASF R3-11 supported Cu oxygen scavenger and molecular sieves (Aldrich, 4Å).

The compounds 6,6-pentamethylenefulvene, $\text{C}_5\text{H}_5\text{CH}_2\text{Ph}$, $(\text{C}_5\text{H}_4\text{C}(\text{=CH}_2)\text{Ph})\text{Li}$, $(\text{C}_5\text{H}_4\text{CMe}_2\text{Ph})\text{TiCl}_3$ (the catalyst used in Example 1), $(\text{C}_5\text{H}_4\text{SiMe}_2\text{Ph})\text{TiCl}_3$ (the catalyst used in Example 3), 10 $(\text{C}_5\text{H}_4\text{CMe}_2\text{-3,5-MeC}_6\text{H}_3)\text{TiCl}_3$ (the catalyst used in Example 2) and $\text{B}(\text{C}_6\text{F}_5)_3$ were prepared according to procedures known as such. 6,6-Diethylfulvene was prepared analogously to 6,6-pentamethylenefulvene from cyclopentadiene and 3-pentanone. $(\text{C}_5\text{H}_4\text{CMe}_2\text{Ph})\text{TiMe}_3$ (used in Examples 10 and 11) was prepared through modification of a known 15 procedure by reaction of $(\text{C}_5\text{H}_4\text{CMe}_2\text{Ph})\text{TiCl}_3$ with either Me_2Mg or MeMgI . The preparations of other titanium complexes are disclosed hereafter in the Preparation Examples A to F.

A toluene solution of MAO (26 wt%, Akzo Nobel Chemicals), MAO supported on silica (5 wt%, Witco) and $[\text{PhNMe}_2\text{H}][\text{B}(\text{C}_6\text{F}_5)_4]$. (Akzo 20 Nobel Chemicals) were used as such.

NMR spectra were recorded on Varian Gemini 200/300 and Unity 500 spectrometers.

The ^1H NMR spectra were referenced to resonances of residual protons in the deuterated solvents. Chemicals shifts (δ) are given 25 relative to tetramethylsilane (downfield shifts are positive). GC analyses were performed on a HP 6890 instrument equipped with a HP-1 dimethylpolysiloxane column (19095 Z-123). GC-MS analyses were conducted using a HP 5973 mass-selective detector attached to a HP 6890 GC instrument. Elemental analyses are the average of a least 30 two independent determinations.

Preparation example A

Preparation of $(\text{C}_5\text{H}_4\text{CH}_2\text{Ph})\text{TiCl}_3$, to be used in Example 4.

a) Preparation of $(\text{C}_5\text{H}_4\text{CH}_2\text{Ph})\text{Li}$

35 To a solution of 11.3 mmol n-BuLi in 30 ml of diethyl ether/hexane at -40°C , 1.87 g (12.0 mmol) of CpHCH_2Ph [2] was added dropwise. The reaction mixture was allowed to warm to room temperature and stirred overnight. The solvents were removed in

vacuo. The white residue was stripped with pentane. After rinsing with 3x 10 ml of pentane and after drying in vacuo 1.55 g (9.6 mmol, 85%) of a white solid was isolated. - ^1H NMR (300 MHz, THF-d_8): δ 8.04-7.92 (m, 4H, Ph o- and m-H), 7.82 (m, 1H, Ph p-H), 6.33 (t, $^3J_{\text{HH}} = 2.6$ Hz, 2H, Cp), 6.29 (t, $^3J_{\text{HH}} = 2.6$ Hz, 2H, Cp), 4.61 (s, 2H, CH_2) - ^{13}C NMR (75.4 MHz, THF-d_8): δ 148.2 (Ph C ipso), 130.5 (Ph o- or m-CH), 129.5 (Ph o- or m-CH), 126.3 (Ph p-CH), 120.0 (Cp C ipso), 105.1 (Cp CH), 103.9 (Cp CH), 39.0 (CH_2)

b) Preparation of $(\text{C}_5\text{H}_4\text{CH}_2\text{Ph})\text{TiCl}_3$

10 To a solution of 1.42 g (8.8 mmol) $(\text{CpCH}_2\text{Ph})\text{Li}$ in 40 ml of methylene chloride, cooled at -40°C , 0.96 ml (1.7 g, 8.9 mmol) titanium(IV) chloride was added. The reaction mixture was stirred at ambient temperatures overnight. The methylene chloride was removed in vacuo and the green-brown residue was stripped with pentane. After
15 extraction with toluene, the extract was evaporated in vacuo and the extract residue was dissolved in methylene chloride. Brown crystals were obtained after cooling to -40°C . Yield: 1.58 g (5.1 mmol, 58%) - ^1H NMR (300 MHz, C_6D_6): δ 7.09-7.03 (m, 3H, Ph m- and p-H), 6.82 (m, 2H, Ph o-H), 5.98 (m, 4H, Cp), 3.71 (s, 2H, CH_2) - ^{13}C NMR (75.4 MHz, C_6D_6): δ 142.9, 138.3 (Ph and Cp C ipso), 128.9, 127.2 (Ph CH, one signal overlapped by solvent), 123.6, 123.2 (Cp CH), 37.7 (CH_2) -
20 Anal. Calcd for $\text{C}_{12}\text{H}_{11}\text{TiCl}_3$: C, 46.57; H, 3.58; Ti, 15.48. Found: C, 47.07; H, 3.75, Ti, 15.38.

25 **Preparation example B**

Preparation of $(\text{C}_5\text{H}_4\text{CET}_2\text{Ph})\text{TiCl}_3$, to be used in Example 5.

a) Preparation of $\text{C}_5\text{H}_4(\text{TMS})\text{CET}_2\text{Ph}$

To a solution of 4.85 g (58 mmol) PhLi in 200 ml of diethyl ether, cooled at -50°C , 8.0 g (60 mmol) of 6,6-diethyl fulvene [1]
30 was added dropwise. The reaction mixture was allowed to warm to room temperature and was stirred for 3 hours. After 3 hours the yellow solution was cooled with an ice bath and 7.6 ml (6.5 g, 60 mmol) of trimethylsilyl chloride was added dropwise. The mixture was allowed to warm to room temperature and stirred overnight. The reaction
35 mixture was poured into 250 ml of ice water. The water layer was extracted with 2x 100 ml of light petroleum, after which the combined organic layers were rinsed with 200 ml of brine. The organic phase was dried on MgSO_4 . After evaporating the low-boiling volatiles in vacuo, the residue was distilled using a Kogelruhr-apparatus. The

product distilled at 110 °C at 0.5 mm Hg as a mixture of isomers.

Yield: 9.21 g (32 mmol, 55%) - ^1H NMR (300 MHz, CDCl_3 , main isomer): δ 7.28 (m, 4H, Ph o- and m-H), 7.18 (m, 1H, Ph p-H), 6.40 (m, 1H, C_5H_4), 6.31 (s, 1H, C_5H_4), 6.22 (m, 1H, C_5H_4), 3.27 (s, 1H, C_5H_4), 2.02 (m, 4H, C- $\underline{\text{CH}_2}$ -CH₃), 0.72 (m, 6H, C-CH₂- $\underline{\text{CH}_3}$), 0.06 (s, 9H, TMS)

b) Preparation of $(\text{C}_5\text{H}_4\text{CEt}_2\text{Ph})\text{TiCl}_3$

To a solution of 6.30 g (22 mmol) of A.1 in 40 ml of methylene chloride, cooled at -40 °C, 2.45 ml (4.2 g, 22 mmol) of titanium chloride was added. The mixture was allowed to warm to room temperature and stirred overnight. The methylene chloride was removed in vacuo and the residue was stripped with pentane. Extraction with methylene chloride and cooling to -60 °C afforded red-brown crystals of the title compound. Yield: 5.63 g (15.3 mmol, 70%) - ^1H NMR (300 MHz, C_6D_6): δ 7.24 (d, $^3J_{\text{HH}} = 7.3$ Hz, 2H, Ph o-H), 7.17 (t, $^3J_{\text{HH}} = 7.3$ Hz, 2H, Ph m-H), 7.06 (t, $^3J_{\text{HH}} = 7.3$ Hz, 1H, Ph p-H), 6.26 (t, $^3J_{\text{HH}} = 2.8$ Hz, 2H, Cp), 6.04 (t, $^3J_{\text{HH}} = 2.8$ Hz, 2H, Cp), 2.06 (m (dq), 2H, C- $\underline{\text{CH}_2}$ -CH₃), 1.86 (m (dq), 2H, C- $\underline{\text{CH}_2}$ -CH₃), 0.51 (t, $^3J_{\text{HH}} = 7.3$ Hz, 6H, C-CH₂- $\underline{\text{CH}_3}$) - ^{13}C NMR (75.4 MHz, C_6D_6): δ 154.8 (Ph C ipso), 142.1 (Cp C ipso), 128.8 (Ph o-CH), 128.3 (Ph m-CH, overlap with solvent), 127.2 (Ph p-CH), 123.1, 121.8 (Cp CH), 48.6 ($\text{C}(\text{C}_2\text{H}_5)_2$ ipso), 29.3 (C- $\underline{\text{CH}_2}$ -CH₃), 8.5 (C-CH₂- $\underline{\text{CH}_3}$) - Anal. Calcd for $\text{C}_{16}\text{H}_{19}\text{TiCl}_3$: C, 52.57; H, 5.24. Found: C, 52.75; H, 5.27.

Preparation example C

25 **Preparation of $\{\text{CpC}[(\text{CH}_2)_5]\text{Ph}\}\text{TiCl}_3$, to be used in Example 6**

a) Preparation of $\text{C}_5\text{H}_4(\text{TMS})\text{C}[(\text{CH}_2)_5]\text{Ph}$

To a solution of 4.00 g (48 mmol) PhLi in 200 ml of diethyl ether, cooled at -50 °C, 6.95 g (48 mmol) of 6,6-pentamethylenefulvene [1] was added dropwise. The reaction mixture was allowed to warm to room temperature and was stirred for 3 hours. After 3 hours the yellow solution was cooled with an ice bath and 6.4 ml (5.5 g, 51 mmol) of trimethylsilyl chloride was added dropwise. The mixture was allowed to warm to room temperature and stirred overnight. The reaction mixture was poured into 250 ml of ice water. The water layer was extracted with 2x 100 ml of light petroleum, after which the combined organic layers were rinsed with 200 ml of brine. The organic phase was dried on MgSO_4 . After evaporating the low-boiling volatiles in vacuo, the residue was distilled using a

Kogelruhr-apparatus. The product distilled at 165 °C at 0.4 torr as a mixture of isomers. Yield: 8.96 g (30 mmol, 63%) - ¹H NMR (300 MHz, CDCl₃, main isomer): δ 7.40 (m, 2H, Ph o-H), 7.33 (m, 2H, Ph m-H), 7.15 (m, 1H, Ph p-H), 6.43 (m, 2H, C₅H₄), 6.15 (s, 1H, C₅H₄), 3.27 (s, 1H, C₅H₄), 2.17 (m, 4H, α-CH₂), 1.65-1.40 (m, 6H, β- and γ-CH₂), - 0.03 (s, 9H, TMS)

b) Preparation of {C₅H₄C[(CH₂)₅]Ph}TiCl₃

Titanium chloride (1.4 ml, 2.4 g, 12.7 mmol) was added to a solution of 3.70 g (12.5 mmol) of C₅H₄(TMS)C[(CH₂)₅]Ph in 40 ml of methylene chloride, cooled at -40 °C. The reaction mixture was allowed to warm to room temperature and was stirred overnight. The methylene chloride was removed in vacuo and the residue was stripped with pentane. The residue was extracted with methylene chloride. Crystallization from a 1:1 mixture of CH₂Cl₂:pentane afforded red-brown crystals of the desired compound in 78% yield (3.68 g, 9.7 mmol). - ¹H NMR (300 MHz, C₆D₆): δ 7.16-7.06 (m, 4H, Ph o- and m-H), 7.01 (m, 1H, Ph p-H), 6.31 (t, ³J_{HH} = 2.8 Hz, 2H, Cp), 5.97 (t, ³J_{HH} = 2.8 Hz, 2H, Cp), 2.45 (d, ²J_{HH} = 13.2 Hz, 2H, α-CH₂ (eq)), 1.88 (m, 2H, α-CH₂ (ax)), 1.37 (br, 3H, β- and γ-CH₂), 1.25-1.05 (m, 3H, β- and γ-CH₂) - ¹³C NMR (75.4 MHz, C₆D₆): δ 156.0 (Ph C ipso), 142.1 (Cp C ipso), 129.2 (Ph o-CH), 127.9 (Ph m-CH), 126.8 (Ph p-CH), 123.2, 120.9 (Cp CH), 45.1 (C[(CH₂)₅] ipso), 35.8 (α-CH₂), 26.1 (γ-CH₂), 22.4 (β-CH₂) - Anal. Calcd for C₁₇H₁₉TiCl₃: C, 54.08; H, 5.07; Ti, 12.69. Found: C, 53.93; H, 4.90; Ti, 12.62.

Preparation example D

Preparation of [C₅H₄C(=CH₂)Ph]TiCl₃, to be used in Example 7

To a solution of 0.61 ml (1.06 g, 5.6 mmol) titanium chloride in 40 ml of methylene chloride, cooled at -50 °C, 1.80 g (5.6 mmol) [C₅H₄C(=CH₂)Ph]Li was added. The reaction mixture was allowed to warm to room temperature and was stirred overnight. The volatiles were removed in vacuo and the green-black residue was stripped with pentane. Extraction with pentane afforded small analytically pure amounts of the desired compound. - ¹H NMR (300 MHz, C₆D₆): δ 7.2-7.05 (m, 5H, Ph), 6.35 (t, ³J_{HH} = 2.7 Hz, 2H, Cp), 6.01 (t, ³J_{HH} = 2.7 Hz, 2H, Cp), 5.58 (s, 1H, =CH₂), 5.20 (s, 1H, =CH₂) - ¹³C NMR (75.4 MHz, C₆D₆): δ 142.5 (Ph C ipso), 139.7 (Cp C ipso), 139.6 (C(=CH₂) ipso), 128.8, 128.7, 128.5 (Ph CH), 123.4, 121.1 (Cp CH), 120.5 (C(=CH₂) -

Anal. Calcd for $C_{13}H_{11}TiCl_3$: C, 48.57; H, 3.45; Ti, 14.90. Found: C, 48.71; H, 3.55; Ti, 14.78.

Preparation Example E

5 Preparation of $C_5H_3(3-SiMe_3)CMe_2Ph]TiCl_3$, to be used in Example 8

a) Preparation of $C_5H_3(SiMe_3)_2CMe_2Ph$

To a solution of 2.25 g (11.8 mmol) $(C_5H_4CMe_2Ph)Li$ [5] in 50 ml of diethyl ether and 20 ml of THF, cooled in ice water, 1.5 ml (1.3 g, 11.9 mmol) $TMSCl$ was added dropwise. The mixture was allowed to
10 warm to room temperature and was stirred overnight. The yellow solution was cooled in ice water and 4.8 ml (12 mmol) of a 2.5M n-BuLi solution in hexanes was added. After warming up to room temperature the mixture was stirred for 4 hours. The white suspension was cooled in ice water and 1.6 ml (1.4 g, 12.7 mmol) $TMSCl$ was added
15 dropwise. The mixture was allowed to warm to room temperature and stirred overnight. The yellow suspension was poured into 125 ml ice water. The water layer was extracted with 50 ml of light petroleum and the combined organic layers were dried on $MgSO_4$. After evaporation of low-boiling volatiles, the residue was distilled using
20 a Kogelruhr-apparatus. The product distilled at 115 °C at 0.8 Torr. Yield: 2.87 g (8.7 mmol, 74%) - 1H NMR (200 MHz, $CDCl_3$): δ 7.4-7.1 (m, 5H, Ph), 6.40 (d, $^3J_{HH} = 2.2$ Hz, 2H, Cp), 6.20 (t, $^3J_{HH} = 2.1$ Hz, 1H, Cp), 1.53 (s, 6H, CMe_2), -0.03 (s, 18H, TMS)

b) Preparation of $[C_5H_3(3-SiMe_3)CMe_2Ph]TiCl_3$

25 To a solution of 0.92 ml (1.6 g, 8.4 mmol) $TiCl_4$ in 50 ml of methylene chloride, cooled at -50 °C, 2.75 g (8.4 mmol) of $C_5H_3(SiMe_3)_2CMe_2Ph$ was added. The reaction mixture was allowed to warm to room temperature and was stirred overnight. The volatiles were removed in vacuo and the residue was stripped with pentane.
30 Extraction with methylene chloride and cooling down to -60 °C afforded 2.76 g (6.7 mmol, 80%) of the desired compound. - 1H NMR (300 MHz, C_6D_6): δ 7.1-6.85 (m, 5H+1H, Ph+Cp), 6.57 (m, 1H, Cp), 6.53 (m, 1H, Cp), 1.63 (s, 6H, CMe_2), 0.12 (s, 9H, TMS) - ^{13}C NMR (75.4 MHz, C_6D_6): δ 158.5 (Ph C ipso), 148.5 (Cp C ipso), 144.1 (Cp C(TMS) ipso), 128.7, 128.6, 126.7, 126.1, 124.6 (Ph+Cp CH), 41.2 (CMe_2 ipso), 29.3, 29.0 (CMe_2), -0.8 (TMS) - Anal. Calcd for $C_{17}H_{23}SiTiCl_3$: C, 49.84; H, 5.66; Ti, 11.69. Found: C, 49.70; H, 5.68; Ti, 11.59.
35

Preparation example F

Preparation of $[C_5H_3(3-SiMe_3)CMe_2-3,5-Me_2C_6H_3]TiCl_3$, to be used in

Example 9

a) Preparation of $C_5H_3(SiMe_3)_2CMe_2-3,5-Me_2C_6H_3$

5 To a solution of 1.15 g (5.3 mmol) $[CpCMe_2-3,5-Me_2C_6H_3]Li$ in 50 ml of diethyl ether, cooled with ice water, 0.7 ml (0.6 g, 5.5 mmol) trimethylsilyl chloride was added dropwise. The reaction mixture was allowed to warm to room temperature and was stirred overnight. The white suspension was cooled to $-30\text{ }^{\circ}C$ and 5.4 mmol of a 2.5M solution

10 of n-BuLi in hexanes was added dropwise. After stirring for 3 hours at ambient temperature, the reaction vessel was placed in ice water and 0.8 ml (0.7 g, 6.4 mmol) trimethylsilyl chloride was added. The reaction mixture was allowed to warm up to room temperature and was stirred overnight. The mixture was poured into 100 ml of ice water.

15 The water layer was extracted twice with 50 ml portions of light petroleum, and the combined organic layers were dried over $MgSO_4$. Kugelrohr-distillation at $160\text{ }^{\circ}C$ and 0.4 Torr yielded 1.26 g (3.5 mmol, 66%) of the title compound - 1H NMR (200 MHz, $CDCl_3$): δ 6.90 (s, 2H, Ar o-H), 6.78 (s, 1H, Ar p-H), 6.37 (m, 2H, Cp), 6.19 (m, 1H,

20 Cp), 2.24 (s, 6H, ArMe), 1.51 (s, 6H, CMe_2), -0.05 (s, 18H, TMS)

b) Preparation of $[C_5H_3(3-SiMe_3)CMe_2-3,5-Me_2C_6H_3]TiCl_3$

To a solution of 0.35 ml (0.6 g, 3.2 mmol) titanium chloride in 40 ml of methylene chloride, cooled at $-40\text{ }^{\circ}C$, 1.18 g (3.3 mmol) of $C_5H_3(SiMe_3)_2CMe_2-3,5-Me_2C_6H_3$ was added dropwise. The reaction mixture

25 was allowed to warm to room temperature and was stirred overnight. The volatiles were removed in vacuo and the residue was stripped with pentane. Extraction with pentane yielded 1.02 g (2.3 mmol, 72%) of light-brown crystals. - 1H NMR (300 MHz, C_6D_6): δ 6.96 (m, 1H, Cp), 6.69 (s, 2H, Ar o-H), 6.64 (m, 1+1H, Cp + p-H), 6.55 (m, 1H, Cp),

30 2.08 (s, 6H, ArMe), 1.70 (s, 6H, CMe_2), 0.13 (s, 9H, TMS) - ^{13}C NMR (75.4 MHz, C_6D_6): δ 159.1 (Ar C ipso), 148.5 (Cp C ipso), 144.1 (Cp C(TMS) ipso), 137.9 (Ar m-C ipso), 128.8, 128.4, 127.8, 124.7 (Cp CH + Ar p-CH), 124.1 (Ar m-H), 41.2 (CMe_2 ipso), 29.3, 29.2 (CMe_2), 21.5 (ArMe), -0.9 (TMS)

Preparation example G**Preparation of $(C_5H_4CMe_2Ph)Ti(CH_2Ph)_3$, to be used in Example 10**

To a stirred solution of 0.52 g of $(C_5H_4CMe_2Ph)TiCl_3$ (1.54 mmol) in 30 ml of diethyl ether, cooled at $-40\text{ }^\circ\text{C}$, a solution of benzyl magnesium bromide (4.62 mmol) in diethyl ether was added dropwise. The mixture was allowed to warm to room temperature and was stirred for 3 hours. The solvent was removed in vacuo. The red solid was extracted with pentane. Cooling to $-40\text{ }^\circ\text{C}$ yielded red crystals of the desired product (560 mg, 1.11 mmol, 72%) - 1H NMR (500 MHz, C_6D_6): δ 7.17-7.11 (m, 10H, Ph *m*- and *o*-H and Bz *m*-H), 7.02 (m, 1H, Ph *p*-H), 6.90 (t, $^3J_{HH} = 7.5$ Hz, 3H, Bz *p*-H), 6.81 (d, $^3J_{HH} = 7.5$ Hz, 6H, Bz *o*-H), 5.74 (ps. t, $^3J_{HH} = 2.8$ Hz, 2H, Cp), 5.50 (ps. t, $^3J_{HH} = 2.8$ Hz, 2H, Cp), 2.97 (s, 6H, Ti- CH_2), 1.38 (s, 6H, CMe_2) - ^{13}C -NMR (125.7 MHz, C_6D_6): δ 149.6 (s, Ph C *ipso*), 149.1 (s, Bz C *ipso*), 146.7 (s, Cp C *ipso*), 128.8 (dd, $^1J_{CH}=158$ Hz, Bz *m*-CH, overlap with solvent), 128.5 (d, $^1J_{CH}=151$ Hz, Ph *m*-CH, overlap with solvent), 127.0 (dm, $^1J_{CH}=161$ Hz, Bz *o*-CH), 126.5 (dm, $^1J_{CH}=156$ Hz, Ph *o*-CH), 126.4 (dm, $^1J_{CH}=156$ Hz, Ph *p*-CH), 123.0 (dt, $^1J_{CH}=160$ Hz, Bz *p*-CH), 118.4 (dm, $^1J_{CH}=168$ Hz, Cp CH), 113.5 (dm, $^1J_{CH}=172$ Hz, Cp CH), 93.5 (t, $^1J_{CH}=123$ Hz, Ti- CH_2), 40.5 (s, CMe_2), 30.2 (q, $^1J_{CH}=122$ Hz, CMe_2) - Anal. Calcd for $C_{35}H_{36}Ti$: C, 83.32; H, 7.19; Ti, 9.49. Found: C, 82.63; H, 7.32; Ti, 9.35.

Example 1: Catalytic ethene conversion with $(C_5H_4CMe_2Ph)TiCl_3/MAO$

The reactions were performed in a stainless steel 1L autoclave (Medimex), fully temperature and pressure controlled and equipped with solvent and catalyst injection systems. In a typical experiment, the autoclave was evacuated and heated for 45 min at $90\text{ }^\circ\text{C}$ prior to use.

The reactor was then brought to the desired temperature, charged with 200 ml of toluene and pressurized with ethene. After equilibrating for 15 min, the appropriate amount of MAO/toluene was injected together with 25 ml of toluene. Subsequently a mixture of 2.50 g cyclooctane (internal standard) and 1.0 ml (0.87 g) of a 15 mM stock solution of the titanium complex in toluene was injected, together with 25 ml of toluene, to start the reaction. During the run the ethene pressure was kept constant to within 0.2 bar of the initial pressure by replenishing flow. After the specified run time, the reactor was vented and the residual MAO was destroyed by addition of 20 ml of ethanol. Samples of the reaction mixture were taken to

analyze and quantify the soluble components. Polymeric product was stirred for 90 min in acidified ethanol and repeatedly rinsed with ethanol and light petroleum on a glass frit. The polymer was initially dried in air and subsequently in vacuo at 70 °C overnight.

5 The results of the catalytic experiments are summarized in Table 1 (ethene conversion with the $(C_5H_4CMe_2Ph)TiCl_3/MAO$ catalyst system) and Table 2 (ethene conversion with the $(C_5H_4CMe_2Ph)TiCl_3/MAO$ catalyst system).

In these experiments, the C_6 fraction consists predominantly of 10 1-hexene (99+ %), with traces of 2- and 3-hexenes. The only detectable product of the C_8 fraction is 1-octene. The C_{10} fraction is a mixture of isomers with either vinyl (90%), vinylidene (5%) or internal olefinic (5%) unsaturation, and consists predominantly of 5-methylnon-1-ene (75-85%). Higher olefins ($C_{12}-C_{24}$) constitute less 15 than 0.5 wt% of the total amount of product formed.

Table 1:

Test nr.	P(ethene) MPa	T °C	C_6 g (wt%)	C_8 g (wt%)	C_{10} g (wt%)	PE g (wt%)	Productivity $kg(C_6) mol(Ti)^{-1} h^{-1}$
1	0.2	30	8.0 (87)	0.1 (0.8)	1.0 (11)	0.2 (1.6)	1066
2	0.5	30	20.9 (83)	0.3 (1.2)	3.5 (14)	0.5 (1.8)	2787
3	1.0	30	47.2 (86)	0.9 (1.6)	5.1 (9)	1.4 (2.6)	6292
4	0.5	50	12.4 (83)	0.2 (1.1)	1.6 (11)	0.7 (4.6)	1653
5	0.5	80	3.3 (76)	0.05(0.9)	0.2 (4)	0.8 (19)	440

20 Toluene solvent, 15 μmol Ti, Al:Ti = 1000, 30 min run time

Table 2:

Test nr.	Run time min	Al:Ti	C_6 g (wt%)	C_8 g (wt%)	C_{10} g (wt%)	PE g (wt%)	Productivity $kg(C_6) mol(Ti)^{-1} h^{-1}$
6	15	1000	16.6 (89)	0.2 (1.0)	1.4 (8)	0.4 (2.2)	4413
7	30	1000	20.9 (83)	0.3 (1.2)	3.5 (14)	0.5 (1.8)	2787
8	60	1000	27.1 (80)	0.4 (1.2)	5.5 (16)	0.9 (2.7)	1809
9	30	500	15.2 (86)	0.2 (1.2)	1.8 (10)	0.5 (2.8)	2029

25

Toluene solvent, 15 μmol Ti, 30 °C, 0.5 MPa ethene

Comparative example A: Catalytic ethene conversion with $(C_5H_4CMe_3)TiCl_3/MAO$

30 The general procedure and conditions of example 1 were followed, using the $(C_5H_4CMe_3)TiCl_3/MAO$ catalyst system. The results of the catalytic experiment are listed in Table 3.

Table 3:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
a	0.5	30	0.5 (17)	0.1 (3)	0.1 (4)	2.4 (76)	72

5 **Example 2: Catalytic ethene conversion with (C₅H₄CMe₂-3,5-Me₂C₆H₃)TiCl₃/MAO**

The general procedure and conditions of example 1 were followed, using the (C₅H₄CMe₂-3,5-Me₂C₆H₃)TiCl₃/MAO catalyst system. The results of the catalytic experiments are listed in Table 4.

10

Table 4:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
10	0.5	30	7.9 (94)	0.02 (0.2)	0.4 (5)	0.1 (1.3)	1052
11	0.5	50	4.5 (93)	0.03 (0.6)	0.2 (4)	0.1 (2.1)	599

15 **Example 3: Catalytic ethene conversion with (C₅H₄SiMe₂Ph)TiCl₃/MAO**

The general procedure and conditions of example 1 were followed, using the (C₅H₄SiMe₂Ph)TiCl₃/MAO catalyst system. The results of the catalytic experiments are listed in Table 5. Higher olefins (C₁₂-C₂₄) are also formed, constituting about 8 wt% of the

20 total amount of product formed.

Table 5:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	C ₁₂ -C ₂₄ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
12	0.5	30	2.1 (36)	0.3 (5)	0.4 (7)	0.5 (8)	2.6 (44)	284
13	0.5	50	2.6 (47)	0.4 (7)	0.4 (7)	0.4 (7)	1.7 (32)	352

25

Example 4: Catalytic ethene conversion with (C₅H₄CH₂Ph)TiCl₃/MAO

The general procedure and conditions of example 1 were followed, using the (C₅H₄CH₂Ph)TiCl₃/MAO catalyst system. The results of the catalytic experiments are listed in Table 6. Higher olefins

30 (C₁₂-C₂₄) are also formed, constituting about 9 wt% of the total amount of products formed.

Table 6:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	C ₁₂ -C ₂₄ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
14	0.5	30	2.7 (42)	0.4 (6)	0.6 (9)	0.6 (9)	2.2 (34)	361
15	0.5	50	3.0 (54)	0.3 (6)	0.5 (9)	0.5 (9)	1.2 (22)	405

5 Example 5: Catalytic ethene conversion with (C₅H₄CET₂Ph)TiCl₃/MAO

The general procedure and conditions of example 1 were followed, using the (C₅H₄CET₂Ph)TiCl₃/MAO catalyst system. The results of the catalytic experiments are listed in Table 7.

10 Table 7:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
16	0.5	30	18.5 (88)	0.05 (0.3)	1.4 (7)	1.0 (4.6)	2462
17	0.5	50	8.7 (84)	0.03 (0.3)	0.6 (5)	1.0 (9.9)	1159

Example 6: Catalytic ethene conversion with {CpC[(CH₂)₅]Ph}TiCl₃/MAO

15 The general procedure and conditions of example 1 were followed, using the {C₅H₄C[(CH₂)₅]Ph}TiCl₃/MAO catalyst system. The results of the catalytic experiments are listed in Table 8.

Table 8:

20

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
18	0.5*	30	24.4 (87)	0.1 (0.3)	2.9 (10)	0.6 (2.0)	3248
19	0.5*	50	12.0 (86)	0.1 (0.4)	1.2 (9)	0.7 (5.2)	1593
20	0.5**	30	16.4 (91)	0.04 (0.2)	1.4 (8)	0.2 (1.3)	4362

* 30 min run time, ** 15 min run time

Example 7: Catalytic ethene conversion with [C₅H₄C(=CH₂)Ph]TiCl₃/MAO

25 The general procedure of example 1 was followed, using the [C₅H₄C(=CH₂)Ph]TiCl₃/MAO catalyst system. The conditions and results of the catalytic experiments are listed in Table 9.

Table 9: Catalytic ethene conversion with the [C₅H₄C(=CH₂)Ph]TiCl₃/MAO

30 system (toluene solvent, 15 μmol Ti, Al:Ti = 1000, 30 min run time)

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
21	0.5	30	17.3 (88)	0.1 (0.3)	1.4 (7)	0.9 (4.7)	2307
22	0.5	50	10.9 (86)	0.03(0.2)	0.7 (6)	1.1 (8.3)	1449

Example 8: Catalytic ethene conversion with $[\text{C}_5\text{H}_3(3\text{-SiMe}_3)\text{CMe}_2\text{Ph}]\text{-TiCl}_3/\text{MAO}$

The general procedure and conditions of example 1 were followed, using the $[\text{C}_5\text{H}_3(3\text{-SiMe}_3)\text{CMe}_2\text{Ph}]\text{TiCl}_3/\text{MAO}$ catalyst system.
5 The results of the catalytic experiments are listed in Table 10.

Table 10:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
23	0.5	30	25.2 (85)	0.6 (2.1)	3.3 (11)	0.4 (1.2)	3357
24	0.5	50	20.1 (84)	0.4 (1.5)	3.3 (14)	0.3 (1.3)	2683
25	0.5	80	8.0 (88)	0.1 (1.4)	0.8 (8)	0.2 (2.1)	1069

10

Example 9: Catalytic ethene conversion with $[\text{C}_5\text{H}_3(3\text{-SiMe}_3)\text{CMe}_2\text{-3,5-Me}_2\text{C}_6\text{H}_3]\text{TiCl}_3/\text{MAO}$

The general procedure and conditions of example 1 were followed, using the $[\text{Cp}(\text{TMS})\text{CMe}_2\text{-3,5-Me}_2\text{C}_6\text{H}_3]\text{TiCl}_3/\text{MAO}$ catalyst
15 system. The results of the catalytic experiments are listed in Table 11. For the run at 30°C, the C₆ fraction consists of 99.9% 1-hexene, and the C₁₀ fraction of 94% 5-methylnon-1-ene.

Table 11:

20

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
26	0.5	30	40.1 (84)	0.1 (0.2)	7.0 (15)	0.3 (0.6)	5347
27	0.5	50	25.7 (82)	0.1 (0.3)	4.8 (15)	0.6 (1.9)	3427

Example 10: Catalytic ethene conversion with $(\text{C}_5\text{H}_4\text{CMe}_2\text{Ph})\text{Ti}(\text{CH}_2\text{Ph})_3/\text{MAO}$

The general procedure and conditions of example 1 were followed, using the $(\text{C}_5\text{H}_4\text{CMe}_2\text{Ph})\text{Ti}(\text{CH}_2\text{Ph})_3/\text{MAO}$ catalyst system. The
25 results of the catalytic experiments are listed in Table 12.

Table 12:

30

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
28	0.5	30	23.8 (82)	0.3 (1.1)	4.6 (16)	0.5 (1.6)	3175
29	0.5	50	18.6 (78)	0.3 (1.1)	4.0 (17)	0.8 (3.3)	2480

Example 11: Catalytic ethene conversion with $(C_5H_4CMe_2Ph)TiMe_3/MAO$

The general procedure and conditions of example 1 were followed, using the $(C_5H_4CMe_2Ph)TiMe_3/MAO$ catalyst system. The results of the catalytic experiment are listed in Table 13.

5

Table 13:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
30	0.5	30	25.7 (81)	0.4 (1.2)	5.2 (16)	0.5 (1.5)	3428
31	0.5	50	18.1 (79)	0.2 (1.0)	3.8 (17)	0.7 (3.2)	2412

10 **Example 12: Catalytic ethene conversion with $(C_5H_4CMe_2Ph)TiMe_3/MAO/SiO_2$**

The reaction was performed in a stainless steel 1 l autoclave (Medimex), fully temperature and pressure controlled and equipped with solvent and catalyst injection systems. Prior to use the autoclave was preheated in vacuo for 45 min at 90 °C. The reactor was cooled to 30 °C, charged with 200 ml of toluene and pressurized with ethene. After equilibrating for 15 min, a slurry of 2.05 g of 5 wt% MAO/SiO₂ in 10 ml of toluene was injected together with 30 ml of toluene. Subsequently a mixture of 2.50 g cyclooctane (internal standard) and 1.0 ml (0.87 g) of a 15 mM stock solution of $(C_5H_4CMe_2Ph)TiMe_3$ in toluene was injected, together with 25 ml of toluene, to start the reaction. During reaction the ethene pressure was kept constant to within 0.2 bar of the initial pressure by replenishing flow. After 30 min the reactor was vented and the remaining residual MAO was destroyed by addition of 20 ml of ethanol. Samples of the reaction mixture were taken to analyze and quantify the soluble components. The solids (polyethene and silica support) were stirred in acidified ethanol for 90 min and rinsed repeatedly with ethanol and light petroleum on a glass frit. The material was dried in air overnight and subsequently in vacuo at 70 °C overnight, yielding 1.7 g of which the polyethene fraction was not determined. The results of the catalytic experiment are listed in Table 14; the conditions were: toluene solvent, 15 μmol Ti, Al:Ti = 250, 30 min run time. Weight percentages calculated on C₆-C₁₀ products only.

35

Table 14:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
32	0.5	30	13.8 (95)	0.1 (0.8)	0.6 (4)	n.d.	1837

5 **Example 13: Catalytic ethene conversion with**
(C₅H₄CMe₂Ph)TiMe₃/[PhNMe₂H][B(C₆F₅)₄]

The reactions were performed in a stainless steel 500 mL autoclave (Medimex), fully temperature and pressure controlled and equipped with solvent and catalyst injection systems. Prior to use
 10 the autoclave was preheated in vacuo for 45 min at 90 °C. The reactor was cooled to the desired temperature, charged with 150 ml of toluene and pressurized with ethene. After equilibrating for 15 min, a suspension of 16.5 µmol [PhNMe₂H][B(C₆F₅)₄] in 5 ml of toluene was injected together with 25 ml of toluene. Subsequently a mixture of
 15 2.50 g cyclooctane (internal standard) and 1.0 ml (0.87 g) of a 15 mM stock solution of the titanium trimethyl complex in toluene was injected, together with 25 ml of toluene, to start the reaction. During reaction the ethene pressure was kept constant to within 0.2 bar of the initial pressure by replenishing flow. After the desired
 20 run time, the reactor was vented and samples of the reaction mixture were taken to analyze and quantify the soluble components. The polymer was repeatedly rinsed with ethanol and light petroleum on a glass frit. The polymer was dried in air overnight and subsequently dried in vacuo at 70 °C overnight. The results of the catalytic
 25 experiments are listed in Table 15. The conditions were: toluene solvent, 15 µmol Ti, B:Ti = 1.1, 30 min run time

Table 15:

Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
33	0.5	30	14.6 (90)	0.2 (1.3)	1.2 (7)	0.3 (2.0)	1948
34	0.5	50	14.0 (82)	0.2 (1.1)	2.2 (13)	0.6 (3.4)	1867

30

Example 14: Catalytic ethene conversion with (C₅H₄CMe₂Ph)TiMe₃/B(C₆F₅)₃

The reaction was performed in a stainless steel 500 mL autoclave (Medimex), fully temperature and pressure controlled and
 35 equipped with solvent and catalyst injection systems. Prior to use the autoclave was preheated in vacuo for 45 min at 90 °C. The reactor

was cooled to the desired temperature, charged with 150 ml of toluene and pressurized with ethene. After equilibrating for 15 min, 1.0 ml (0.87 g) of a 16.5 mM stock solution of $B(C_6F_5)_3$ in toluene was injected together with 25 ml of toluene. Subsequently a mixture of 5 2.50 g cyclooctane (internal standard) and 1.0 ml (0.87 g) of a 15 mM stock solution of the titanium trimethyl complex in toluene was injected, together with 25 ml of toluene, to start the reaction. During reaction the ethene pressure was kept constant to within 0.2 bar of the initial pressure by replenishing flow. After the desired 10 run time, the reactor was vented and samples of the reaction mixture were taken to analyze and quantify the soluble components. The polymer was repeatedly rinsed with ethanol and light petroleum on a glass frit. The polymer was dried in air overnight and subsequently dried in vacuo at 70 °C overnight. The results of the catalytic 15 experiment are listed in Table 16. The conditions were: toluene solvent, 15 μ mol Ti, B:Ti = 1.1, 30 min run time.

Table 16:

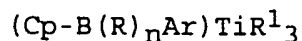
Test nr.	P(ethene) MPa	T °C	C ₆ g (wt%)	C ₈ g (wt%)	C ₁₀ g (wt%)	PE g (wt%)	Productivity kg(C ₆) mol(Ti) ⁻¹ h ⁻¹
35	0.5	30	5.8 (88)	0.1 (0.9)	0.3 (5)	0.4 (6.6)	776

20

C L A I M S

1. A catalyst system for the selective trimerization of olefins comprising a transition metal complex characterized in that said catalyst comprises

- a) a half-sandwich substituted cyclopentadienyl titanium complex
of formula



wherein

Cp is a cyclopentadienyl type ligand, optionally substituted,

B is a bridging group, based on a single atom selected from the groups 13 to 16 inclusive of the Periodic System,

Ar is a aromatic group, optionally substituted,

R is, independently, hydrogen, or a hydrocarbon residue, optionally being substituted and optionally containing heteroatoms, or groups R and B are joined together to form a ring,

n is an integer equal to the (valency of B minus 2), and

R¹ is a mono-anionic group, and

- b) an activator.

2. A catalyst system according to claim 1 wherein said group B consists of carbon or silicon, Ar is phenyl, optionally substituted or being part of a larger aromatic entity,

R¹ is a halide, or mono-anionic hydrocarbon residue optionally containing heteroatoms, and

n is 2, then R is a mono-anionic hydrocarbon residue, optionally containing heteroatoms, or

n is 1, then R is a di-anionic hydrocarbon residue, optionally containing heteroatoms.

3. A catalyst system according to claims 1 or 2 in which Ar is a phenyl group, substituted or not at the meta-or para-position,

B is based on a carbon atom,

n is 2, then groups R are, independently, methyl, or ethyl; or

n is 1, then group R is =CH₂, or forms when R is C₄H₈ or C₅H₁₀ together with group B a dianionic cyclic group

Cp is C_5H_4 or $C_5H_3(SiMe_3)$, and

R^1 is chlorine, methyl, or benzyl.

4. A catalyst system according to any of the claims 1 to 3
5 wherein said catalyst complex is supported on a carrier.

5. A catalyst system according to any of the claims 1 to 4
wherein said activator is methylalumoxane, a salt of a non-
coordinating anion, or a Lewis acid capable of abstracting an anion
10 from said transition metal complex.

6. A catalyst system according to claims 4 and 5 wherein the
activator is methylalumoxane and the molar ratio of Ti:Al is from
1:100 to 1:1000.
15

7. A catalyst system according to any of the claims 1 to 6,
wherein said catalyst system further comprises a scavenger.

8. A process to trimerize olefinic compounds, which comprises
20 carrying out said trimerization in the presence of a catalyst system
according to any of the preceding claims under trimerization
conditions.

9. A process according to claim 8, wherein said olefin is
25 selected from C_2 - C_{20} olefins or mixtures of two or more of these
olefins.

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07C2/32 B01J31/22

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07C B01J

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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A	EP 0 608 447 A (PHILLIPS PETROLEUM CO) 3 August 1994 (1994-08-03) cited in the application claims 20-24	8,9
X	SASSMANNSHAUSEN J ET AL: "HALF-SANDWICH COMPLEXES OF TITANIUM AND ZIRCONIUM WITH PENDANT PHENYL SUBSTITUENTS. THE INFLUENCE OF ANSA-ARYL COORDINATION ON THE POLYMERISATION ACTIVITY OF HALF-SANDWICH CATALYSTS" JOURNAL OF ORGANOMETALLIC CHEMISTRY, ELSEVIER-SEQUOIA S.A. LAUSANNE, CH, vol. 592, 1999, pages 84-94, XP001033815 ISSN: 0022-328X cited in the application table 1	1-7
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☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

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P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

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Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

G document member of the same patent family

Date of the actual completion of the international search

12 November 2001

Date of mailing of the international search report

21/11/2001

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

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